

PS Claim 10; Col 151; 127pp; English.
 XX
 CC The present invention describes isolated DNA (I) encoding at least one
 CC osteogenically active region of human osteogenic protein-1 in prepro form
 CC (OPI-PP), murine OPI-PP, murine OP2-PP or human OP2-PP. Also described
 CC are: (A) DNA related to (I) encoding a polypeptide able to form dimers
 CC that can induce cartilage and endochondral bone formation in a mammal
 CC when implanted in a matrix; (B) vectors containing (I) or related DNA;
 CC (C) host cells transformed with this vector; (D) DNA (I') encoding a
 CC prepro- or pro-OPI, and related vectors and transformed cells; (E)
 CC osteogenic protein (II) produced by expression of transformed mammalian
 CC cells, able to induce bone and cartilage formation; (F) mature OPI
 CC secreted from mammalian cells following expression of the sequence that
 CC encodes hOP1-PP; and (G) production of an active osteogenic composition
 CC by truncating mature OPI protein. Host cells of (C) are used to produce
 CC proteins able to induce cartilage and bone formation, e.g. for correction
 CC of acquired or congenital craniofacial defects or other skeletal or
 CC dental disorders; to heal non-union fractures; to repair cartilage, e.g.
 CC in osteoarthritis, or generally wherever bone formation is required. The
 CC proteins induce complete development of endochondral bone, including
 CC vascularisation, mineralisation and bone marrow differentiation. The
 CC present sequence represents a human OPI fragment. (Updated on 20-MAR-2003
 XX to correct PA field.)

SQ Sequence 97 AA;

Query Match 100.0%; Score 111; DB 2; Length 97;
 Best Local Similarity 100.0%; Pred. No. 4.3e-07;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 INPSETVPKPCCAPTQINAI 20
 DB 52 INPSETVPKPCCAPTQINAI 71

RESULT 4

ID AAW95444 standard; protein; 97 AA.

AC AAW95444;

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DT 26-MAR-1999 (first entry)

XX

DE Conserved 6 cysteine skeleton fragment from human OPI.

XX

KW Cystic kidney disease; renal; therapeutic; osteogenic protein; OP;

KW bone morphogenic protein; BMP; growth factor-beta superfamily;

KW polycystic kidney disease; multicystic dysplastic kidney disease;

KW uremic medullary cystic disease; human.

XX

OS Homo sapiens.

XX

PN WO985061-A1.

XX

PD 12-NOV-1998.

XX

PF 06-MAY-1998; 98WO-US009268.

XX

PR 07-MAY-1997; 97US-0045909P.

XX

PA (BIOL) BIOPEN INC.

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PI Gjorstrup P, Harris R;

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DR WPI; 1999-070084/06.

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PT Treating cystic kidney disease - using renal therapeutic agents or
 PT sequences encoding them, especially from the osteogenic protein/bone
 PT morphogenic protein family.

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PS Claim 3; Page 5-6; 67pp; English.

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CC The invention relates to methods for treating cystic kidney diseases. The

method comprises administering an effective amount of a renal therapeutic agent or a polynucleotide encoding the therapeutic agent. The therapeutic agent is preferably a soluble or membrane bound polypeptide, e.g. a member of the osteogenic protein/bone morphogenic protein (OP/BMP) family within a transforming growth factor-beta superfamily of proteins. It is especially one of the polypeptides hOP1, hOP1-PP, OP1-18aa, OPS, OP7, OP1-16Ser, OP1-16Leu, OP1-16Met, OP1-16Ala, OP1-16Val, mOP1, mOP1-PP, hOP2, hOP2-PP, hOP2-Ala, hOP2-Pro, hOP2-Arg, or hOP2-Ser or their biologically active homologues. The method is used to treat humans having, or at risk of, cystic kidney disease, e.g. autosomal recessive (infantile) polycystic disease, multicystic dysplastic kidney disease, uremic medullary cystic disease, and autosomal dominant polycystic kidney disease. The present sequence represents a human osteogenic protein 1 (OPI) species defining the active region in the conserved 6 cysteine skeleton in the active region.

SQ Sequence 97 AA;

Query Match 100.0%; Score 111; DB 2; Length 97;
 Best Local Similarity 100.0%; Pred. No. 4.3e-07;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 INPSETVPKPCCAPTQINAI 20
 DB 52 INPSETVPKPCCAPTQINAI 71

RESULT 5

ID AAP95681 standard; protein; 98 AA.

AC AAP95681;

XX

DT 25-MAR-2003 (revised)

DT 21-AUG-1990 (first entry)

XX

DE Human osteogenic protein 1 (OP1-I) for osteogenic device.

XX

KW Osteogenic device; osteogenic protein; endochondral bone;

KW biodegradable matrix.

XX

OS Synthetic.

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PN WO8909788-A.

XX

PD 19-OCT-1989.

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PF 08-APR-1988; 88US-00179406.

XX

PR 15-AUG-1988; 88US-00232630.

XX

PR 23-FEB-1989; 88US-0031542.

XX

PR 07-APR-1989; 88WO-US001469.

XX

PA (CREA-) CREATIVE BIOMOLECULES INC.

XX

PI Oppermann H, Kuberaamp T, Rueger D;

XX

DR WPI; 1989-324203/44.

XX

PT Osteogenic devices comprising matrix contg. osteogenic proteins - prep'd
 PT by recombinant techniques.

XX

PS Claim 9; Page 48; 69pp; English.

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CC The protein is capable of inducing endochondral bone formation in
 CC association with a biocompatible, in vivo biodegradable matrix. The
 CC protein is produced by expression of the recombinant DNA in a host cell
 CC and comprises more than one polypeptide chain, with an amino acid
 CC sequence sufficiently duplicative of COP5, COP7, COP16 or OPI. The
 CC protein and the implantable devices enable optimal predictable bone
 CC formation. Clinical applications include correction of acquired and
 CC congenital craniofacial and other skeletal or dental anomalies, induction

CC of local endochondral bone formation in non-union fractures, periodontal
 CC apptns. requiring bone formation and cartilage repair, eg in the
 CC treatment of osteoarthritis. See also AAP95679-PR5692 and AN9097.
 CC (Updated on 25-MAR-2003 to correct PR field.) (Updated on 25-MAR-2003 to
 CC correct PA field.)

SQ Sequence 98 AA:

Query Match 100.0%; Score 111; DB 1; Length 98;
 Best Local Similarity 100.0%; Pred. No. 4.4e-07; Mismatches 0; Indels 0; Gaps 0;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 INPETVPKCCAPTONAIS 20
 Db 53 INPETVPKCCAPTONAIS 72

RESULT 5
 AAP95682
 ID AAP95682 standard; protein; 102 AA.
 XX
 AC AAP95682;
 XX
 DT 25-MAR-2003 (revised)
 DT 21-AUG-1990 (first entry)
 DE Human osteogenic protein 1(OPI-II) for osteogenic device.
 XX
 KW Osteogenic device; osteogenic protein; endochondral bone;
 KW biodegradable matrix.
 XX
 OS Synthetic.
 XX
 PN WO909788-A.
 XX
 PD 19-OCT-1989.
 XX
 PR 08-APR-1988; 88US-00179406.
 XX
 PR 08-APR-1988; 88US-00179406.
 PR 15-AUG-1988; 88US-0032630.
 PR 23-FEB-1989; 89US-00315342.
 PR 07-APR-1989; 89WO-US001469.
 XX
 PA (CREA-) CREATIVE BIOMOLECULES INC.
 XX
 PI Oppermann H, Kuberasamp T, Rueger D;
 DR WPI, 1989-324203/44.
 XX
 PT Osteogenic devices comprising matrix contg. osteogenic proteins - prepa.
 PT by recombinant techniques.
 XX
 PS Claim 10; Page 49; 69pp; English.

XX
 CC The Protein is capable of inducing endochondral bone formation in
 CC association with a biocompatible, in vivo biodegradable matrix. The
 CC protein is produced by expression of the recombinant DNA in a host cell
 CC and comprises more than one polypeptide chain, with an amino acid
 CC sequence sufficiently duplicative of COP5, COP7, COP16 or OPI. The
 CC protein and the implantable devices enable optimal predictable bone
 CC formation. Clinical applications include correction of acquired and
 CC congenital craniofacial and other skeletal or dental anomalies, induction
 CC of local endochondral bone formation in non-union fractures, Periodontal
 CC applicns. requiring bone formation and cartilage repair, eg in the
 CC treatment of osteoarthritis. See also AAP95679-PR5692 and AN9097.
 CC (Updated on 25-MAR-2003 to correct PR field.) (Updated on 25-MAR-2003 to
 CC correct PA field.)
 XX
 SQ Sequence 102 AA:

Query Match 100.0%; Score 111; DB 1; Length 102;
 Best Local Similarity 100.0%; Pred. No. 4.5e-07; Mismatches 0; Indels 0; Gaps 0;

RESULT 7
 AAR53360
 ID AAR53360 standard; protein; 102 AA.
 XX
 AC AAR53360;
 XX
 DT 25-MAR-2003 (revised)
 DT 01-JUL-2002 (revised)
 DT 06-JUN-1994 (first entry)
 DE Osteogenic protein OP7.
 XX
 KW Osteogenic protein; bone; cartilage; matrix; osteoarthritis; repair;
 KW vascularisation; mineralisation; differentiation.
 XX
 OS Homo sapiens.
 XX
 PN US5266683-A.
 XX
 PD 30-NOV-1993.
 XX
 PR 21-FEB-1992; 92US-00841646.
 PR 08-APR-1988; 88US-00179406.
 PR 15-AUG-1988; 88US-00232630.
 PR 23-FEB-1989; 89US-00315342.
 PR 17-OCT-1989; 89US-00422633.
 PR 17-OCT-1989; 89US-00422639.
 PR 22-FEB-1990; 90US-00483933.
 PR 20-AUG-1990; 90US-00569950.
 PR 07-SEP-1990; 90US-00579465.
 PR 18-OCT-1990; 90US-00599543.
 PR 18-OCT-1990; 90US-00599543.
 PR 21-NOV-1990; 90US-00616374.
 PR 04-DEC-1990; 90US-00621849.
 PR 04-DEC-1990; 90US-00621888.
 PR 22-FEB-1991; 91US-00660162.
 PR 20-DEC-1991; 91US-00810560.
 PR 28-JAN-1992; 92US-00827052.
 XX
 PA (SYTC) STRYKER CORP.
 XX
 PT Kuberasampath T, Ozkaynak E, Rueger DC, Pang RHL, Oppermann H;
 XX
 DR WPI; 1993-395405/49.
 N-PSDB; AA053141.
 XX
 PT New pure mammalian osteogenic proteins - induce cartilage and
 PT endochondral bone formation when in association with a matrix.
 XX
 PS Claim 7; Col 69-72; 128pp; English.

XX
 CC This sequence is a fragment of the osteogenic protein OPI and is
 CC designated OP7. The sequence is a 102 C-terminal region and functional
 CC domain of OPI. The osteogenic protein when in association with a matrix
 CC can induce at the locus of an implant the full development cascade of
 CC endochondral bone formation including vascularisation, mineralisation and
 CC bone marrow differentiation. The osteogenic protein can also be used to
 CC repair both bone and cartilage in the treatment of osteoarthritis
 CC (Updated on 01-JUN-2002 to add missing PA field.) (Updated on 25-MAR-2003
 CC to correct PF field.) (Updated on 25-MAR-2003 to correct PR field.)
 XX
 SQ Sequence 102 AA:

Query Match 100.0%; Score 111; DB 2; Length 102;
 Best Local Similarity 100.0%; Pred. No. 4.5e-07;

RESULT 154
 US-09-019-339B-2
 Sequence 2, Application US/09019339B
 Patent No. 6201195
 386 INPETVPKPCCAPTQNAIS 405

RESULT 154

US-09-019-339B-2

Sequence 2, Application US/09019339B

GENERAL INFORMATION:

APPLICANT: RUGGER, David C

APPLICANT: TUCKER, Marjorie M

TITLE OF INVENTION: MATRIX-FREE OSTEOGENIC DEVICES, IMPLANTS AND

TITLE OF INVENTION: METHODS OF USE THEREOF

NUMBER OF SEQUENCE: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: James F. Haley, Jr., Esq. c/o FISH & NEAVE

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STATE: New York

COUNTRY: United States of America

ZIP: 10020

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30 (EPO)

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/019,339B

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APPLICATION NUMBER:

FILING DATE:

CLASSIFICATION:

ATTORNEY/AGENT INFORMATION:

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REFERENCE/DOCKET NUMBER: CPP-147

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INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 431 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-09-019-339B-2

Query Match Score 111; DB 3; Length 431;
 Best Local Similarity 100.0%; Pred. No. 2.3e-07;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy	1	INPETVPKPCCAPTQNAIS	20
Db	386	INPETVPKPCCAPTQNAIS	405

RESULT 155